

**ASSEMBLY BILL**

**No. 465**

**Introduced by Assembly Member Cogdill**

February 16, 2005

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An act to amend Sections 11100, 11100.1, 11106, and 11107.1 of the Health and Safety Code, relating to controlled substances.

LEGISLATIVE COUNSEL'S DIGEST

AB 465, as introduced, Cogdill. Controlled substances: iodine.

(1) Existing law generally provides that any manufacturer, wholesaler, retailer, or other person or entity in this state that sells, transfers, or otherwise furnishes to any person or entity in this or any other state any of a list of substances shall submit a report to the Department of Justice of all of those transactions, and shall submit an application to, and obtain a permit for the conduct of that business from, the Department of Justice, as specified. Any person who does not submit a report as required, who submits a false report, or who sells, transfers, or furnishes a substance without a permit is guilty of a crime, punishable as specified.

Existing law does not include iodine in the list of substances for which a report must be provided, or a permit to conduct business required, but existing law does make it a misdemeanor for any person to sell or purchase more than 8 ounces of iodine in any 30-day period, other than tincture of iodine, any topical solution containing iodine that is equal to or less than \$100, or iodine sold to specified licensed entities that sell, transfer, or furnish the iodine to a licensed pharmacy, physician, dentist, podiatrist, or veterinarian.

This bill would delete the provision prohibiting the sale or purchase of 8 ounces of iodine in any 30-day period. This bill would instead add iodine and tincture of iodine to the list of substances with respect

to which transactions must be reported and for which a permit to conduct business must be obtained, except in specified circumstances. By increasing the scope of persons to whom existing crimes are applicable, this bill would impose a state-mandated local program upon local government.

(2) Existing law provides that the reporting requirement is not applicable to any manufacturer or wholesaler licensed by the California State Board of Pharmacy that sells, transfers, or otherwise furnishes a substance to specified entities, provided the manufacturer or wholesaler submits records of any suspicious sales or transfers as determined by the Department of Justice.

This bill would provide that the reporting requirement is also not applicable to a state-licensed health care facility that administers or furnishes a substance to its patients, or to the sale, transfer, furnishing, or receipt of any betadine or povidone solution with an iodine content not exceeding 1% in containers of 8 ounces or less, or any tincture of iodine not exceeding 2% in containers of one ounce or less, that is sold over the counter.

(3) Existing law provides that the permit requirement is not applicable to specified entities, including retailers and other persons, that are licensed by the California State Board of Pharmacy and also registered with the federal Drug Enforcement Agency.

This bill would remove retailers and other persons from this exemption. It would provide that the permit requirement is also not applicable to any pharmacist or other authorized person who sells or furnishes a substance upon the prescription of a physician, dentist, podiatrist, or veterinarian; or to any state-licensed health care facility, physician, dentist, podiatrist, veterinarian, or veterinary food-animal drug retailer that administers or furnishes a substance to a patient. The bill would add an exemption from the permit requirement for the sale, transfer, furnishing, or receipt of any betadine or povidone solution with an iodine content not exceeding 1% in containers of 8 ounces or less, or any tincture of iodine not exceeding 2% in containers of one ounce or less, that is sold over the counter.

(4) The California Constitution requires the state to reimburse local agencies and school districts for certain costs mandated by the state. Statutory provisions establish procedures for making that reimbursement.

This bill would provide that no reimbursement is required by this act for a specified reason.

Vote: majority. Appropriation: no. Fiscal committee: yes.  
State-mandated local program: yes.

*The people of the State of California do enact as follows:*

- 1 SECTION 1. Section 11100 of the Health and Safety Code is
- 2 amended to read:
- 3 11100. (a) Any manufacturer, wholesaler, retailer, or other
- 4 person or entity in this state that sells, transfers, or otherwise
- 5 furnishes any of the following substances to any person or entity
- 6 in this state or any other state shall submit a report to the
- 7 Department of Justice of all of those transactions:
- 8 (1) Phenyl-2-propanone.
- 9 (2) Methamphetamine.
- 10 (3) Ethylamine.
- 11 (4) D-lysergic acid.
- 12 (5) Ergotamine tartrate.
- 13 (6) Diethyl malonate.
- 14 (7) Malonic acid.
- 15 (8) Ethyl malonate.
- 16 (9) Barbituric acid.
- 17 (10) Piperidine.
- 18 (11) N-acetylanthranilic acid.
- 19 (12) Pyrrolidine.
- 20 (13) Phenylacetic acid.
- 21 (14) Anthranilic acid.
- 22 (15) Morpholine.
- 23 (16) Ephedrine.
- 24 (17) Pseudoephedrine.
- 25 (18) Norpseudoephedrine.
- 26 (19) Phenylpropanolamine.
- 27 (20) Propionic anhydride.
- 28 (21) Isosafrole.
- 29 (22) Safrole.
- 30 (23) Piperonal.
- 31 (24) Thionylchloride.
- 32 (25) Benzyl cyanide.
- 33 (26) Ergonovine maleate.
- 34 (27) N-methylephedrine.
- 35 (28) N-ethylephedrine.

- 1 (29) N-methylpseudoephedrine.
- 2 (30) N-ethylpseudoephedrine.
- 3 (31) Chloroephedrine.
- 4 (32) Chloropseudoephedrine.
- 5 (33) Hydriodic acid.
- 6 (34) Gamma-butyrolactone, including butyrolactone;
- 7 butyrolactone gamma; 4-butyrolactone; 2(3H)-furanone dihydro;
- 8 dihydro-2(3H)-furanone; tetrahydro-2-furanone; 1,2-butanolide;
- 9 1,4-butanolide; 4-butanolide; gamma-hydroxybutyric acid
- 10 lactone; 3-hydroxybutyric acid lactone and 4-hydroxybutanoic
- 11 acid lactone with Chemical Abstract Service number (96-48-0).
- 12 (35) 1,4-butanediol, including butanediol; butane-1,4-diol;
- 13 1,4-butylene glycol; butylene glycol; 1,4-dihydroxybutane;
- 14 1,4-tetramethylene glycol; tetramethylene glycol; tetramethylene
- 15 1,4-diol with Chemical Abstract Service number (110-63-4).
- 16 (36) Red phosphorous, including white phosphorous,
- 17 hypophosphorous acid and its salts, ammonium hypophosphite,
- 18 calcium hypophosphite, iron hypophosphite, potassium
- 19 hypophosphite, manganese hypophosphite, magnesium
- 20 hypophosphite, and sodium hypophosphite.
- 21 (37) *Iodine or tincture of iodine.*
- 22 (38) Any of the substances listed by the Department of Justice
- 23 in regulations promulgated pursuant to subdivision (b).
- 24 (b) The Department of Justice may adopt rules and regulations
- 25 in accordance with Chapter 3.5 (commencing with Section
- 26 11340) of Part 1 of Division 3 of Title 2 of the Government Code
- 27 that add substances to subdivision (a) if the substance is a
- 28 precursor to a controlled substance and delete substances from
- 29 subdivision (a). However, no regulation adding or deleting a
- 30 substance shall have any effect beyond March 1 of the year
- 31 following the calendar year during which the regulation was
- 32 adopted.
- 33 (c) (1) (A) Any manufacturer, wholesaler, retailer, or other
- 34 person or entity in this state, prior to selling, transferring, or
- 35 otherwise furnishing any substance specified in subdivision (a) to
- 36 any person or business entity in this state or any other state, shall
- 37 require (A) a letter of authorization from that person or business
- 38 entity that includes the currently valid business license number or
- 39 federal Drug Enforcement Administration (DEA) registration
- 40 number, the address of the business, and a full description of how

the substance is to be used, and (B) proper identification from the purchaser. The manufacturer, wholesaler, retailer, or other person or entity in this state shall retain this information in a readily available manner for three years. The requirement for a full description of how the substance is to be used does not require the person or business entity to reveal their chemical processes that are typically considered trade secrets and proprietary information.

(B) For the purposes of this paragraph, “proper identification” for in-state or out-of-state purchasers includes two or more of the following: federal tax identification number; seller’s permit identification number; city or county business license number; license issued by the California Department of Health Services; registration number issued by the Federal Drug Enforcement Administration; precursor business permit number issued by the Bureau of Narcotic Enforcement of the California Department of Justice; ~~motor vehicle operator’s~~ *driver’s* license; or other identification issued by a state.

(2) (A) Any manufacturer, wholesaler, retailer, or other person or entity in this state that exports a substance specified in subdivision (a) to any person or business entity located in a foreign country shall, on or before the date of exportation, submit to the Department of Justice a notification of that transaction, which notification shall include the name and quantity of the substance to be exported and the name, address, and, if assigned by the foreign country or subdivision thereof, business identification number of the person or business entity located in a foreign country importing the substance.

(B) The department may authorize the submission of the notification on a monthly basis with respect to repeated, regular transactions between an exporter and an importer involving a substance specified in subdivision (a), if the department determines that a pattern of regular supply of the substance exists between the exporter and importer and that the importer has established a record of utilization of the substance for lawful purposes.

(d) (1) Any manufacturer, wholesaler, retailer, or other person or entity in this state that sells, transfers, or otherwise furnishes a substance specified in subdivision (a) to a person or business entity in this state or any other state shall, not less than 21 days

1 prior to delivery of the substance, submit a report of the  
2 transaction, which includes the identification information  
3 specified in subdivision (c), to the Department of Justice. The  
4 Department of Justice may authorize the submission of the  
5 reports on a monthly basis with respect to repeated, regular  
6 transactions between the furnisher and the recipient involving the  
7 substance or substances if the Department of Justice determines  
8 that a pattern of regular supply of the substance or substances  
9 exists between the manufacturer, wholesaler, retailer, or other  
10 person or entity that sells, transfers, or otherwise furnishes the  
11 substance or substances and the recipient of the substance or  
12 substances, and the recipient has established a record of  
13 utilization of the substance or substances for lawful purposes.

14 (2) The person selling, transferring, or otherwise furnishing  
15 any substance specified in subdivision (a) shall affix his or her  
16 signature or otherwise identify himself or herself as a witness to  
17 the identification of the purchaser or purchasing individual, and  
18 shall, if a common carrier is used, maintain a manifest of the  
19 delivery to the purchaser for three years.

20 (e) This section shall not apply to any of the following:

21 (1) Any pharmacist or other authorized person who sells or  
22 furnishes a substance upon the prescription of a physician,  
23 dentist, podiatrist, or veterinarian.

24 (2) Any physician, dentist, podiatrist, or veterinarian who  
25 administers or furnishes a substance to his or her patients.

26 (3) Any manufacturer or wholesaler licensed by the California  
27 State Board of Pharmacy that sells, transfers, or otherwise  
28 furnishes a substance to a licensed pharmacy, physician, dentist,  
29 podiatrist, veterinarian, or retail distributor as defined in  
30 subdivision (h), provided that the manufacturer or wholesaler  
31 submits records of any suspicious sales or transfers as determined  
32 by the Department of Justice.

33 (4) Any analytical research facility that is registered with the  
34 federal Drug Enforcement Administration of the United States  
35 Department of Justice.

36 (5) *A state-licensed health care facility that administers or*  
37 *furnishes a substance to its patients.*

38 (6) (A) Any sale, transfer, furnishing, or receipt of any  
39 product that contains ephedrine, pseudoephedrine,  
40 norpseudoephedrine, or phenylpropanolamine and which is

lawfully sold, transferred, or furnished over the counter without a prescription pursuant to the federal Food, Drug, and Cosmetic Act (21 U.S.C. Sec. 301 et seq.) or regulations adopted thereunder. However, this section shall apply to preparations in solid or liquid dosage form, except pediatric liquid forms, as defined, containing ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine where the individual transaction involves more than three packages or nine grams of ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine.

(B) Any ephedrine, pseudoephedrine, norpseudoephedrine, or phenylpropanolamine product subsequently removed from exemption pursuant to Section 814 of Title 21 of the United States Code shall similarly no longer be exempt from any state reporting or permitting requirement, unless otherwise reinstated pursuant to subdivision (d) or (e) of Section 814 of Title 21 of the United States Code as an exempt product.

~~(6)~~

*(7) The sale, transfer, furnishing, or receipt of any betadine or povidone solution with an iodine content not exceeding 1 percent in containers of eight ounces or less, or any tincture of iodine not exceeding 2 percent in containers of one ounce or less, that is sold over the counter.*

(8) Any transfer of a substance specified in subdivision (a) for purposes of lawful disposal as waste.

(f) (1) Any person specified in subdivision (a) or (d) who does not submit a report as required by that subdivision or who knowingly submits a report with false or fictitious information shall be punished by imprisonment in a county jail not exceeding six months, by a fine not exceeding five thousand dollars (\$5,000), or by both the fine and imprisonment.

(2) Any person specified in subdivision (a) or (d) who has previously been convicted of a violation of paragraph (1) shall, upon a subsequent conviction thereof, be punished by imprisonment in the state prison, or by imprisonment in a county jail not exceeding one year, by a fine not exceeding one hundred thousand dollars (\$100,000), or by both the fine and imprisonment.

(g) (1) Except as otherwise provided in subparagraph (A) of paragraph ~~(5)~~ (6) of subdivision (e), it is unlawful for any

1 manufacturer, wholesaler, retailer, or other person to sell,  
2 transfer, or otherwise furnish a substance specified in subdivision  
3 (a) to a person under 18 years of age.

4 (2) Except as otherwise provided in subparagraph (A) of  
5 paragraph ~~(5)~~ (6) of subdivision (e), it is unlawful for any person  
6 under 18 years of age to possess a substance specified in  
7 subdivision (a).

8 (3) Notwithstanding any other law, it is unlawful for any retail  
9 distributor to (i) sell in a single transaction more than three  
10 packages of a product that he or she knows to contain ephedrine,  
11 pseudoephedrine, norpseudoephedrine, or phenylpropanolamine,  
12 or (ii) knowingly sell more than nine grams of ephedrine,  
13 pseudoephedrine, norpseudoephedrine, or phenylpropanolamine,  
14 other than pediatric liquids as defined. Except as otherwise  
15 provided in this section, the three package per transaction  
16 limitation or nine gram per transaction limitation imposed by this  
17 paragraph shall apply to any product that is lawfully sold,  
18 transferred, or furnished over the counter without a prescription  
19 pursuant to the federal Food, Drug, and Cosmetic Act (21 U.S.C.  
20 Sec. 301 et seq.), or regulations adopted thereunder, unless  
21 exempted from the requirements of the federal Controlled  
22 Substances Act by the federal Drug Enforcement Administration  
23 pursuant to Section 814 of Title 21 of the United States Code.

24 (4) (A) A first violation of this subdivision is a misdemeanor.

25 (B) Any person who has previously been convicted of a  
26 violation of this subdivision shall, upon a subsequent conviction  
27 thereof, be punished by imprisonment in a county jail not  
28 exceeding one year, by a fine not exceeding ten thousand dollars  
29 (\$10,000), or by both the fine and imprisonment.

30 (h) For the purposes of this article, the following terms have  
31 the following meanings:

32 (1) “Drug store” is any entity described in Code 5912 of the  
33 Standard Industrial Classification (SIC) Manual published by the  
34 United States Office of Management and Budget, 1987 edition.

35 (2) “General merchandise store” is any entity described in  
36 Codes 5311 to 5399, inclusive, and Code 5499 of the Standard  
37 Industrial Classification (SIC) Manual published by the United  
38 States Office of Management and Budget, 1987 edition.



1 (3) “Grocery store” is any entity described in Code 5411 of the  
2 Standard Industrial Classification (SIC) Manual published by the  
3 United States Office of Management and Budget, 1987 edition.

4 (4) “Pediatric liquid” means a nonencapsulated liquid whose  
5 unit measure according to product labeling is stated in  
6 milligrams, ounces, or other similar measure. In no instance shall  
7 the dosage units exceed 15 milligrams of phenylpropanolamine  
8 or pseudoephedrine per five milliliters of liquid product, except  
9 for liquid products primarily intended for administration to  
10 children under two years of age for which the recommended  
11 dosage unit does not exceed two milliliters and the total package  
12 content does not exceed one fluid ounce.

13 (5) “Retail distributor” means a grocery store, general  
14 merchandise store, drugstore, or other related entity, the activities  
15 of which, as a distributor of ephedrine, pseudoephedrine,  
16 norpseudoephedrine, or phenylpropanolamine products, are  
17 limited exclusively to the sale of ephedrine, pseudoephedrine,  
18 norpseudoephedrine, or phenylpropanolamine products for  
19 personal use both in number of sales and volume of sales, either  
20 directly to walk-in customers or in face-to-face transactions by  
21 direct sales. “Retail distributor” includes an entity that makes a  
22 direct sale, but does not include the parent company of that entity  
23 if the company is not involved in direct sales regulated by this  
24 article.

25 (6) “Sale for personal use” means the sale in a single  
26 transaction to an individual customer for a legitimate medical use  
27 of a product containing ephedrine, pseudoephedrine,  
28 norpseudoephedrine, or phenylpropanolamine in dosages at or  
29 below that specified in paragraph (3) of subdivision (g). “Sale for  
30 personal use” also includes the sale of those products to  
31 employers to be dispensed to employees from first-aid kits or  
32 medicine chests.

33 (i) It is the intent of the Legislature that this section shall  
34 preempt all local ordinances or regulations governing the sale by  
35 a retail distributor of over-the-counter products containing  
36 ephedrine, pseudoephedrine, norpseudoephedrine, or  
37 phenylpropanolamine.

38 SEC. 2. Section 11100.1 of the Health and Safety Code is  
39 amended to read:

11100.1. (a) Any manufacturer, wholesaler, retailer, or other person or entity in this state that obtains from a source outside of this state any substance specified in subdivision (a) of Section 11100 shall submit a report of that transaction to the Department of Justice 21 days in advance of obtaining the substance. However, the Department of Justice may authorize the submission of reports within 72 hours, or within a timeframe and in a manner acceptable to the Department of Justice, after the actual physical obtaining of a specified substance with respect to repeated transactions between a furnisher and an obtainer involving the substances, if the Department of Justice determines that the obtainer has established a record of utilization of the substances for lawful purposes. This section does not apply to any person whose prescribing or dispensing activities are subject to the reporting requirements set forth in Section 11164, ~~or to;~~ any manufacturer, ~~or wholesaler, retailer, or other person~~ who is licensed by the California State Board of Pharmacy and also registered with the federal Drug Enforcement Administration of the United States Department of Justice, ~~or to;~~ any analytical research facility that is registered with the federal Drug Enforcement Administration of the United States Department of Justice; *or any state-licensed health care facility.*

(b) (1) Any person specified in subdivision (a) who does not submit a report as required by that subdivision shall be punished by imprisonment in a county jail not exceeding six months, by a fine not exceeding five thousand dollars (\$5,000), or by both that fine and imprisonment.

(2) Any person specified in subdivision (a) who has been previously convicted of a violation of subdivision (a) who subsequently does not submit a report as required by subdivision (a) shall be punished by imprisonment in the state prison, or by imprisonment in a county jail not exceeding one year, by a fine not exceeding one hundred thousand dollars (\$100,000), or by both that fine and imprisonment.

SEC. 3. Section 11106 of the Health and Safety Code is amended to read:

11106. (a) (1) (A) Any manufacturer, wholesaler, retailer, or any other person or entity in this state that sells, transfers, or otherwise furnishes any substance specified in subdivision (a) of Section 11100 to a person or business entity in this state or any

1 other state or who obtains from a source outside of the state any  
2 substance specified in subdivision (a) of Section 11100 shall  
3 submit an application to, and obtain a permit for the conduct of  
4 that business from, the Department of Justice. For any substance  
5 added to the list set forth in subdivision (a) of Section 11100 on  
6 or after January 1, 2002, the Department of Justice may postpone  
7 the effective date of the requirement for a permit for a period not  
8 to exceed six months from the listing date of the substance.

9 (B) An intracompany transfer does not require a permit if the  
10 transferor is a permittee. Transfers between company partners or  
11 between a company and an analytical laboratory do not require a  
12 permit if the transferor is a permittee and a report as to the nature  
13 and extent of the transfer is made to the Department of Justice  
14 pursuant to Section 11100 or 11100.1.

15 (C) This paragraph shall not apply to any manufacturer; *or*  
16 ~~wholesaler, retailer, or other person or entity that~~ *who* is licensed  
17 by the California State Board of Pharmacy and also registered  
18 with the federal Drug Enforcement Administration of the United  
19 States Department of Justice, ~~or to;~~ *any pharmacist or other*  
20 *authorized person who sells or furnishes a substance upon the*  
21 *prescription of a physician, dentist, podiatrist, or veterinarian;*  
22 *any state-licensed health care facility, physician, dentist,*  
23 *podiatrist, veterinarian, or veterinary food-animal drug retailer*  
24 *licensed by the California State Board of Pharmacy that*  
25 *administers or furnishes a substance to a patient; or any*  
26 analytical research facility that is registered with the federal Drug  
27 Enforcement Administration of the United States Department of  
28 Justice.

29 (D) *This paragraph shall not apply to the sale, transfer,*  
30 *furnishing, or receipt of any betadine or povidone solution with*  
31 *an iodine content not exceeding 1 percent in containers of eight*  
32 *ounces or less, or any tincture of iodine not exceeding 2 percent*  
33 *in containers of one ounce or less, that is sold over the counter.*

34 (2) Except as provided in paragraph (3), no permit shall be  
35 required of any manufacturer, wholesaler, retailer, or other  
36 person or entity for the sale, transfer, furnishing, or obtaining of  
37 any product which contains ephedrine, pseudoephedrine,  
38 norpseudoephedrine, or phenylpropanolamine and which is  
39 lawfully sold, transferred, or furnished over the counter without a  
40 prescription or by a prescription pursuant to the federal Food,

1 Drug, and Cosmetic Act (21 U.S.C. Sec. 301 et seq.) or  
2 regulations adopted thereunder.

3 (3) A permit shall be required for the sale, transfer, furnishing,  
4 or obtaining of preparations in solid or liquid dosage form  
5 containing ephedrine, pseudoephedrine, norpseudoephedrine, or  
6 phenylpropanolamine, unless (A) the transaction involves the  
7 sale of ephedrine, pseudoephedrine, norpseudoephedrine, or  
8 phenylpropanolamine products by retail distributors as defined  
9 by this article over the counter and without a prescription, or (B)  
10 the transaction is made by a person or business entity exempted  
11 from the permitting requirements of this subdivision under  
12 paragraph (1).

13 (b) (1) The department shall provide application forms, which  
14 are to be completed under penalty of perjury, in order to obtain  
15 information relating to the identity of any applicant applying for  
16 a permit, including, but not limited to, the business name of the  
17 applicant or the individual name, and if a corporate entity, the  
18 names of its board of directors, the business in which the  
19 applicant is engaged, the business address of the applicant, a full  
20 description of any substance to be sold, transferred, or otherwise  
21 furnished or to be obtained, the specific purpose for the use, sale,  
22 or transfer of those substances specified in subdivision (a) of  
23 Section 11100, the training, experience, or education relating to  
24 this use, and any additional information requested by the  
25 department relating to possible grounds for denial as set forth in  
26 this section, or by applicable regulations adopted by the  
27 department.

28 (2) The requirement for the specific purpose for the use, sale,  
29 or transfer of those substances specified in subdivision (a) of  
30 Section 11100 does not require applicants or permittees to reveal  
31 their chemical processes that are typically considered trade  
32 secrets and proprietary business information.

33 (c) Applicants and permittees shall authorize the department,  
34 or any of its duly authorized representatives, as a condition of  
35 being permitted, to make any examination of the books and  
36 records of any applicant, permittee, or other person, or visit and  
37 inspect the business premises of any applicant or permittee  
38 during normal business hours, as deemed necessary to enforce  
39 this chapter.

1 (d) An application may be denied, or a permit may be revoked  
2 or suspended, for reasons which include, but are not limited to,  
3 the following:

4 (1) Materially falsifying an application for a permit or an  
5 application for the renewal of a permit.

6 (2) If any individual owner, manager, agent, representative, or  
7 employee for the applicant who has direct access, management,  
8 or control for any substance listed under subdivision (a) of  
9 Section 11100, is or has been convicted of a misdemeanor or  
10 felony relating to any of the substances listed under subdivision  
11 (a) of Section 11100, any misdemeanor drug-related offense, or  
12 any felony under the laws of this state or the United States.

13 (3) Failure to maintain effective controls against the diversion  
14 of precursors to unauthorized persons or entities.

15 (4) Failure to comply with this article or any regulations of the  
16 department adopted thereunder.

17 (5) Failure to provide the department, or any duly authorized  
18 federal or state official, with access to any place for which a  
19 permit has been issued, or for which an application for a permit  
20 has been submitted, in the course of conducting a site  
21 investigation, inspection, or audit; or failure to promptly produce  
22 for the official conducting the site investigation, inspection, or  
23 audit any book, record, or document requested by the official.

24 (6) Failure to provide adequate documentation of a legitimate  
25 business purpose involving the applicant's or permittee's use of  
26 any substance listed in subdivision (a) of Section 11100.

27 (7) Commission of any act which would demonstrate actual or  
28 potential unfitness to hold a permit in light of the public safety  
29 and welfare, which act is substantially related to the  
30 qualifications, functions, or duties of a permitholder.

31 (8) If any individual owner, manager, agent, representative, or  
32 employee for the applicant who has direct access, management,  
33 or control for any substance listed under subdivision (a) of  
34 Section 11100, willfully violates or has been convicted of  
35 violating, any federal, state, or local criminal statute, rule, or  
36 ordinance regulating the manufacture, maintenance, disposal,  
37 sale, transfer, or furnishing of any of those substances.

38 (e) Notwithstanding any other provision of law, an  
39 investigation of an individual applicant's qualifications, or the  
40 qualifications of an applicant's owner, manager, agent,

1 representative, or employee who has direct access, management,  
2 or control of any substance listed under subdivision (a) of  
3 Section 11100, for a permit may include review of his or her  
4 summary criminal history information pursuant to Sections  
5 11105 and 13300 of the Penal Code, including, but not limited to,  
6 records of convictions, regardless of whether those convictions  
7 have been expunged pursuant to Section 1203.4 of the Penal  
8 Code, and any arrests pending adjudication.

9 (f) The department may retain jurisdiction of a canceled or  
10 expired permit in order to proceed with any investigation or  
11 disciplinary action relating to a permittee.

12 (g) The department may grant permits on forms prescribed by  
13 it, which shall be effective for not more than one year from the  
14 date of issuance and which shall not be transferable. Applications  
15 and permits shall be uniform throughout the state, on forms  
16 prescribed by the department.

17 (h) Each applicant shall pay at the time of filing an application  
18 for a permit a fee determined by the department which shall not  
19 exceed the application processing costs of the department.

20 (i) A permit granted pursuant to this article may be renewed  
21 one year from the date of issuance, and annually thereafter,  
22 following the timely filing of a complete renewal application  
23 with all supporting documents, the payment of a permit renewal  
24 fee not to exceed the application processing costs of the  
25 department, and a review of the application by the department.

26 (j) Selling, transferring, or otherwise furnishing or obtaining  
27 any substance specified in subdivision (a) of Section 11100  
28 without a permit is a misdemeanor or a felony.

29 (k) (1) No person under 18 years of age shall be eligible for a  
30 permit under this section.

31 (2) No business for which a permit has been issued shall  
32 employ a person under 18 years of age in the capacity of a  
33 manager, agent, or representative.

34 (l) (1) An applicant, or an applicant's employees who have  
35 direct access, management, or control of any substance listed  
36 under subdivision (a) of Section 11100, for an initial permit shall  
37 submit with the application ~~two sets~~ *one set* of 10-print  
38 fingerprint cards for each individual acting in the capacity of an  
39 owner, manager, agent, or representative for the applicant, unless  
40 the applicant's employees are exempted from this requirement by

the Department of Justice. These exemptions may only be obtained upon the written request of the applicant.

(2) In the event of subsequent changes in ownership, management, or employment, the permittee shall notify the department in writing within 15 calendar days of the changes, and shall submit one set of 10-print fingerprint cards for each individual not previously fingerprinted under this section.

SEC. 4. Section 11107.1 of the Health and Safety Code is amended to read:

11107.1. (a) Any manufacturer, wholesaler, retailer, or other person or entity in this state that sells to any person or entity in this state or any other state any quantity of sodium cyanide, potassium cyanide, cyclohexanone, bromobenzene, magnesium turnings, mercuric chloride, sodium metal, lead acetate, palladium black, ~~iodine~~, hydrogen chloride gas, trichlorofluoromethane (fluorotrichloromethane), dichlorodifluoromethane, 1,1,2-trichloro-1,2,2-trifluoroethane (trichlorotrifluoroethane), sodium acetate, or acetic anhydride shall do the following:

(1) (A) Notwithstanding any other provision of law, in any face-to-face or will-call sale, the seller shall prepare a bill of sale which identifies the date of sale, cost of sale, method of payment, the specific items and quantities purchased and the proper purchaser identification information, all of which shall be entered onto the bill of sale or a legible copy of the bill of sale, and shall also affix on the bill of sale his or her signature as witness to the purchase and identification of the purchaser.

(B) For the purposes of this paragraph, “proper purchaser identification” includes a valid ~~motorvehicle operator’s~~ *driver’s* license or other official and valid state-issued identification of the purchaser that contains a photograph of the purchaser, and includes the residential or mailing address of the purchaser, other than a post office box number, the motor vehicle license number of the motor vehicle used by the purchaser at the time of purchase, a description of how the substance is to be used, the Environmental Protection Agency certification number or resale tax identification number assigned to the individual or business entity for which the individual is purchasing any chlorofluorocarbon product, and the signature of the purchaser.

(C) The seller shall retain the original bill of sale containing the purchaser identification information for five years in a readily

1 presentable manner, and present the bill of sale containing the  
2 purchaser identification information upon demand by any law  
3 enforcement officer or authorized representative of the Attorney  
4 General. Copies of these bills of sale obtained by representatives  
5 of the Attorney General shall be maintained by the Department  
6 of Justice for a period of not less than five years.

7 (2) (A) Notwithstanding any other law, in all sales other than  
8 face-to-face or will-call sales the seller shall maintain for a  
9 period of five years the following sales information: the name  
10 and address of the purchaser, date of sale, product description,  
11 cost of product, method of payment, method of delivery, delivery  
12 address, and valid identifying information.

13 (B) For the purposes of this paragraph, “valid identifying  
14 information” includes two or more of the following: federal tax  
15 identification number; resale tax identification number; city or  
16 county business license number; license issued by the State  
17 Department of Health Services; registration number issued by the  
18 federal Drug Enforcement Administration; precursor business  
19 permit number issued by the Bureau of Narcotic Enforcement of  
20 the Department of Justice; ~~motor vehicle operator’s~~ *driver’s*  
21 license; or other identification issued by a state.

22 (C) The seller shall, upon the request of any law enforcement  
23 officer or any authorized representative of the Attorney General,  
24 produce a report or record of sale containing the information in a  
25 readily presentable manner.

26 (D) If a common carrier is used, the seller shall maintain a  
27 manifest regarding the delivery in a readily presentable manner  
28 for a period of five years.

29 (b) Any manufacturer, wholesaler, retailer, or other person or  
30 entity in this state that purchases any item listed in subdivision  
31 (a) of Section 11107.1 shall do the following:

32 (1) Provide on the record of purchase information on the  
33 source of the items purchased, the date of purchase, a description  
34 of the specific items, the quantities of each item purchased, and  
35 the cost of the items purchased.

36 (2) Retain the record of purchase for three years in a readily  
37 presentable manner and present the record of purchase upon  
38 demand to any law enforcement officer or authorized  
39 representative of the Attorney General.



1     ~~(e) (1) Except as provided in paragraph (2), no manufacturer,~~  
2     ~~wholesaler, retailer, or other person or entity shall sell to any~~  
3     ~~individual, and no individual shall buy, more than eight ounces~~  
4     ~~of iodine in any 30-day period.~~

5     ~~(2) For purposes of this section, these requirements do not~~  
6     ~~apply to either of the following:~~

7     ~~(A) Any sale of tincture of iodine or any topical solution~~  
8     ~~containing iodine that is equal to or less than one hundred dollars~~  
9     ~~(\$100).~~

10    ~~(B) Any sale of iodine made to a licensed health care facility,~~  
11    ~~any manufacturer licensed by the State Department of Health~~  
12    ~~Services, or wholesaler licensed by the California State Board of~~  
13    ~~Pharmacy who sells, transfers, or otherwise furnishes the iodine~~  
14    ~~to a licensed pharmacy, physician, dentist, podiatrist, or~~  
15    ~~veterinarian.~~

16    ~~(d)~~

17    ~~(c) (1) A first violation of this section is a misdemeanor.~~

18    ~~(2) Any person who has previously been convicted of a~~  
19    ~~violation of this section shall, upon a subsequent conviction~~  
20    ~~thereof, be punished by imprisonment in a county jail not~~  
21    ~~exceeding one year, by a fine not exceeding one hundred~~  
22    ~~thousand dollars (\$100,000), or both the fine and imprisonment.~~

23    SEC. 5. No reimbursement is required by this act pursuant to  
24    Section 6 of Article XIII B of the California Constitution because  
25    the only costs that may be incurred by a local agency or school  
26    district will be incurred because this act creates a new crime or  
27    infraction, eliminates a crime or infraction, or changes the  
28    penalty for a crime or infraction, within the meaning of Section  
29    17556 of the Government Code, or changes the definition of a  
30    crime within the meaning of Section 6 of Article XIII B of the  
31    California Constitution.